

OCT 25 2004

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**510(K) SUMMARY**

**Given® Diagnostic System with PillCam™ ESO Capsule**

**510(k) Number K041149\_\_\_\_\_**

**Applicant's Name:**

Given Imaging Ltd.  
13 HaYetzira St.  
P.O. Box 258  
New Industrial Zone  
Yokneam 20692, Israel  
Tel.: 011-972-4-9097730  
Fax: 011-972-4-9592466

**Contact Person:**

Shosh Friedman, RAC  
Corporate V.P. Regulatory & Medical Affairs  
Tel: 011-972-4- 9097784  
Fax: 011-972-4-9938060  
Email: shosh@givenimaging.com

**Trade Name:**

Given® Diagnostic System

**Classification Name:**

Ingestible Telemetric Gastrointestinal Capsule Imaging System

**Classification:**

FDA has classified Ingestible Telemetric Gastrointestinal Capsule Imaging System as class II devices (product code 78NZE) and they are reviewed by the Gastroenterology Panel.

**Predicate Device:**

Given® Diagnostic System (K010312, K020341, K022362, K022980, K031033, K032405, and K040248.)

**Performance Standards and Special Controls:**

The Given® Diagnostic System complies with the requirements presented in “Class II Special Controls Guidance Document; Ingestible Telemetric Gastrointestinal Capsule Imaging System; Final Guidance for Industry and FDA” issued on November 28, 2001

**Intended Use:**

The Given® Diagnostic System with the PillCam™ ESO Capsule is intended for the visualization of esophageal mucosa.

**Device Description:**

The Given® Diagnostic System with the PillCam™ ESO Capsule is comprised of three subsystems: PillCam™ ESO Capsule, Data Recorder Set, and RAPID® Workstation.

The PillCam™ ESO Capsule is a wireless, disposable capsule designed to glide smoothly through the esophagus. During its passage, the capsule transmits digital data that is captured by receiving antennas attached to the patient's body. The images are stored in the DataRecorder that is connected to the receiving antennas and is worn on a belt around the waist of the patient. When the test is over, the antennas and recorder are removed from the patient's body. The images from the recorder are downloaded to the RAPID® Workstation for processing and viewing by the physician.

**Substantial Equivalence:**

Given Imaging Ltd. believes that, based on the information provided in this submission, the Given® Diagnostic System with PillCam™ ESO Capsule is substantially equivalent to the market-cleared Given® Diagnostic System (with the M2A® SB Capsule) without raising any new safety and/or efficacy issue.



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

Shosh Friedman, RAC  
Corporate Vice President  
Regulatory & Medical Affairs  
Given Imaging Ltd.  
New Industrial Park  
P.O. Box 258, Yoqneam  
20692 ISRAEL

OCT 25 2004

Re: K041149

Trade/Device Name: Given<sup>®</sup> Diagnostic System with PillCam<sup>™</sup> ESO Capsule  
Regulation Number: 21 CFR §876.1300  
Regulation Name: Ingestible telemetric gastrointestinal capsule imaging system  
Regulatory Class: II  
Product Code: 78 NSI  
Dated: September 12, 2004  
Received: September 22, 2004

Dear Ms. Friedman:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Office of Compliance at one of the following numbers, based on the regulation number at the top of this letter:

21 CFR 876.xxxx	(Gastroenterology/Renal/Urology)	240-276-0115
21 CFR 884.xxxx	(Obstetrics/Gynecology)	240-276-0115
21 CFR 892.xxxx	(Radiology)	240-276-0120
Other		240-276-0100

Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR 807.97). You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its Internet address <http://www.fda.gov/cdrh/dsma/dsmamain.html>

Sincerely yours,



Nancy C. Brogdon  
Director, Division of Reproductive,  
Abdominal, and Radiological Devices  
Office of Device Evaluation  
Center for Devices and Radiological Health

Enclosure

## INDICATIONS FOR USE STATEMENT

**510(k) Number (if known):** K041149

**Device Name:**

Given® Diagnostic System with PillCam™ ESO Capsule

**Indications for Use:**

The Given® Diagnostic System with the PillCam™ ESO Capsule is intended for the visualization of the esophagus.

(PLEASE DO NOT WRITE BELOW THIS LINE -CONTINUE ON ANOTHER PAGE IF NEEDED)

510(k) Number K041149

Prescription Use ✓  
(Per 21 CFR 801.109)

OR

Over the Counter Use \_\_\_\_\_

Nancy C Brozdon  
(Division Sign-Off)  
Division of Reproductive, Abdominal,  
and Radiological Devices  
510(k) Number K041149